

RECEIVED

PTO-1442 REPRODUCED

INFORMATION DISCLOSURE CITATION
IN AN APPLICATION

MAY 29 2001

May 23, 2001

(Use several sheets if necessary)

ATTORNEY DOCKET NO.
3028.1000-000APPLICATION NO.
09/590,211APPLICANT
Guy A. Rouleau and Bernard BraisFILING DATE
June 8, 2000GROUP
1632JUN 01 2001
FBI-CENTER 1600/2900

U.S. PATENT DOCUMENTS

EXAM- INER INI- TIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUB- CLASS	FILING DATE IF APPROPRIATE
AA						
AB						

FOREIGN PATENT DOCUMENTS

	DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB- CLASS	TRANSLATION YES NO
AL	WO98/31800	23 JUL 98	PCT			
AM	WO99/29896	17 JUN 99	PCT			
AN						
AO						

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

AR	Akarsu, A.N., et al., "Genomic Structure of HOXD13 Gene: A Nine Polyalanine Duplication Causes Synpolydactyly in Two Unrelated Families," <i>Human Molecular Genetics</i> , 5(7): 945-952 (1996).
AS	Bienroth, S, et al., "Assembly of a Processive Messenger RNA Polyadenylation Complex," <i>The EMBO Journal</i> , 12(2): 585-594 (1993).
AT	Brais, B., et al., "Using the Full Power of Linkage Analysis in 11 French Canadian Families to Fine Map the Oculopharyngeal Muscular Dystrophy Gene," <i>Neuromuscular Disorder</i> 7(1):S70-S74 (1997).
AU	Brais, B, et al., "The Oculopharyngeal Muscular Dystrophy Locus Maps to the Region of the Cardiac α and β Myosin Heavy Chain Genes on Chromosome 14q11.2-q13," <i>Human Molecular Genetics</i> , 4(3): 429-434 (1995).
AV	Davies, S.W., "Formation of Neuronal Intranuclear Inclusions Underlies the Neurological Dysfunction in Mice Transgenic for the HD Mutation," <i>Cell</i> , 90:537-548 (1997).
AW	DiFiglia, M, et al., "Aggregation of Huntingtin in Neuronal Intranuclear Inclusions and Dystrophic Neurites in Brain," <i>Science</i> , 277: 1990-1993 (1997).
AX	Evans, G.A, et al., "High Efficiency Vectors for Cosmid Microcloning and Genomic Analysis," <i>Gene</i> , 79:9-20 (1989).
AY	Forood, B., et al., "Formation of an Extremely Stable Polyalanine β -Sheet Macromolecule," <i>Biochem. And Biophysical Res. Communications</i> , 211(1): 7-13 (1995).
AZ	Krause, S., et al., "Immunodetection of Poly(A) Binding Protein II in the Cell Nucleus," <i>Experimental Cell Res.</i> , 214: 75-82 (1994).

EXAMINER

DATE CONSIDERED

Joe Wailar

7/8/04

PTO-1449 REPRODUCED

ATTORNEY DOCKET NO.
3028.1000-000APPLICATION NO.
09/590,211INFORMATION DISCLOSURE CITATION
IN AN APPLICATION

May 23, 2001

(Use several sheets if necessary)

APPLICANT
Guy A. Rouleau and Bernard BraisFILING DATE
June 8, 2000GROUP
1632

U.S. PATENT DOCUMENTS

EXAM- INER INI- TIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUB- CLASS	FILING DATE IF APPROPRIATE

FOREIGN PATENT DOCUMENTS

DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB- CLASS	TRANSLATION YES NO

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

AR2	Mundlos, S., et al., "Mutations Involving the Transcription Factor CBFA1 Cause Cleidocranial Dysplasia," <i>Cell</i> , 89: 773-779 (1997).
AS2	Editorials, "DNA-Triplet Repeats and Neurologic Disease," <i>The New England Journal of Med.</i> , 335(16): 1222-1224 (1996).
AT2	Scherzinger, E., et al., "Huntingtin-Encoded Polyglutamine Expansions Form Amyloid-Like Protein Aggregates In Vitro and In Vivo," <i>Cell</i> 90: 549-558 (1997).
AU2	Tome, M.S., et al., "Nuclear Inclusions in Oculopharyngeal Dystrophy," <i>Act Neuropathol.</i> 49: 85-87 (1980).
AV2	Muragaki, Y., et al., "Polyalanine Expansion in Synpolydactyly Might Result from Unequal Crossing-Over of HOXD13," <i>Science</i> 275: 406
AW2	Wells, R.D., "Molecular Basis of Genetic Instability of Triplet Repeats," <i>The Journal of Biological Chem.</i> 271(6): 2875-2878 (1996).
AX2	Wahle, E., et al., "Mammalian Poly(A)-Binding Protein II," <i>J. of Biological Chem.</i> , 268(4): 2937-2945 (1993).
AY2	Wahle E., "A Novel Poly(A)-Binding Protein Acts As a Specificity Factor in the Second Phase of Messenger RNA Polyadenylation," <i>Cell</i> , 66: 759-768 (1991).
AZ2	Nemeth, A., et al., "Isolation of Genomic and cDNA Clones Encoding Bovine Poly(A) Binding Protein II," <i>Nucleic Acids Res.</i> , 23(20): 4034-4041 (1995).
AR3	Riggins, G.J., et al. "Human Genes Containing Polymorphic Trinucleotide Repeats," <i>Nat Genet</i> , 2(3):186-191 (1992).
AS3	Brais, B., et al., "Short GCG Expansions in the PABP2 Gene Cause Oculopharyngeal Muscular Dystrophy," <i>Nature Genetics</i> 18: 164-167 (1998).

EXAMINER

DATE CONSIDERED



7/8/04

TECH CENTER 1600/2800

JUN 01 2001

RECEIVED

PTO-1449 REPRODUCED

**INFORMATION DISCLOSURE CITATION
IN AN APPLICATION**

MAY 29 2001

(Use several sheets if necessary)

ATTORNEY DOCKET NO.
3028.1000-000

APPLICATION NO.
09/590,211

APPLICANT
Guy A. Rouleau and Bernard Brais

FILING DATE
June 8, 2000

GROUP
1632

U.S. PATENT DOCUMENTS

EXAM- INER INI- TIAL	DOCUMENT NUMBER	DATE	NAME	CLASS	SUB- CLASS	FILING DATE IF APPROPRIATE

FOREIGN PATENT DOCUMENTS

DOCUMENT NUMBER	DATE	COUNTRY	CLASS	SUB- CLASS	TRANSLATION YES NO

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)

AT3 ✓	Lamartine, J. et al., "Cloning Sequencing and Chromosomal Assignment of a New cDNA Clone to Xq12-q13 and 14q11," EMBL Data Base Accession Number U12206 (1995).
AU3 ✓	"National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index," EMBL Data Base Accession Number AA618589 (1997).
AV3 ✓	Sullivan, T.B.T., et al., "Oculopharyngeal Muscular Dystrophy (OPMD) - Report and Genetic Studies of an Australian Kindred," <i>Clinical Genetics</i> , 51: 52-55 (1997).
AW3 ✓	Bouchard, J.P. et al., "A Simple Test for the Detection of Dysphagia in Members of Families with Oculopharyngeal Muscular Dystrophy (OPMD)," <i>Can. J. Neurol. Sci.</i> 19(2):296-297 (1992).

EXAMINER

DATE CONSIDERED


